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A method for treating or preventing gastritis in a subject, comprising administering to said subject a therapeutically effective amount of an amylin or an amylin agonist wherein said amylin agonist is not a calcitonin.

2. A method for treating or preventing gastric ulceration in a subject, comprising administering to said subject a therapeutically effective amount of an amylin or an amylin agonist, wherein said amylin agonist is not a calcitonin.

A method of treating or preventing pain, fever, inflammation, arthritis, hypercoagulability, or other condition for which a non-steroidal anti-inflammatory agent would be indicated, comprising administering to subject a therapeutically effective amount of an amylin or an amylin agonist, wherein said amylin agonist is not a calcitonin, and a therapeutically effective amount of a non-steroidal anti-inflammatory agent.

A method of enhancing the analgesic activity of a non-steroidal anti-inflammatory drug in a subject, comprising administering an amylin or an amylin agonist along with said non-steroidal anti-inflammatory drug, wherein said amylin agonist is not a calcitonin.

- 5. The method according to any of claims 1-4, wherein said subject is human.
- 6. The mathod according to any of claims 1-4, wherein said amylin or amylin agonist is administered by a route selected form the group consisting of nasal, oral, pulmonary, transdermal, and buccal administration.
- 7. The method according to any of claims 1-4 wherein said amylin agonist is selected from the group consisting

[PATENTS\224042\224042.APP]

15

of  $^{18}$ Arg $^{25}$  $^{28}$ Pro-h-amylin, des- $^{1}$ Lys $^{18}$ Arg $^{25,28}$ Pro-h-amylin,  $^{18}$ Arg $^{25-}$ <sup>28,29</sup>Pro-h-amylin, des-<sup>1</sup>Lys<sup>18</sup>Arg<sup>25,28,29</sup>Pro-h-amylin, <sup>25,28-29</sup>Pro-hamylin, des-1Lys<sup>25,28,29</sup>Pro-h-amylin, <sup>25</sup>Pro<sup>26</sup>Val<sup>28,29</sup>Pro-h-amylin, <sup>23</sup>Leu<sup>25</sup>Pro<sup>26</sup>Val<sup>28,29</sup>Pro-h-amylin, <sup>23</sup>Leu<sup>25</sup>Pro<sup>26</sup>Val<sup>28</sup>Pro-h-amylin, des-1Lys<sup>23</sup>Leu<sup>25</sup>Pro<sup>26</sup>Val<sup>28</sup>Pro-h-amylin, <sup>18</sup>Arg<sup>23</sup>Leu<sup>25</sup>Pro<sup>26</sup>Val<sup>28</sup>Proh-amylin, 18 Arg23 Leu25,28,29 Pro-h-amylin, 18 Arg23 Leu25,28 Pro-hamylin, <sup>17</sup>Ile Leu<sup>25,28,29</sup>Pro-h-amylin, <sup>17</sup>Ile<sup>25,28,29</sup>Pro-h-amylin, des-Lys<sup>17</sup>Ile<sup>23</sup>Leu<sup>25,28,29</sup>Pro-h-amylin, <sup>17</sup>Ile<sup>18</sup>Arg<sup>23</sup>Leu-h-amylin, <sup>17</sup>Ile<sup>18</sup>Arg<sup>23</sup>Leu<sup>26</sup>∜al<sup>29</sup>Pro-h-amylin, <sup>17</sup>Ile<sup>18</sup>Arg<sup>23</sup>Leu<sup>25</sup>Pro<sup>26</sup>Val<sup>28,29</sup>Pro-h-amylin, 13Thr<sup>21</sup>His<sup>23</sup>Leu<sup>26</sup>Ala<sup>28</sup>Leu<sup>29</sup>Pro<sup>31</sup>Asp-h-amylin, <sup>13</sup>Thr<sup>21</sup>His<sup>23</sup>Leu<sup>26</sup>Ala<sup>29</sup>Pro<sup>31</sup>Asp-h-amylin, des-<sup>1</sup>Lys<sup>13</sup>Thr<sup>21</sup>His<sup>23</sup>Leu<sup>26</sup>Ala<sup>28</sup>Pro<sup>31</sup>Asp-h-amylin, <sup>13</sup>Thr<sup>18</sup>Arg<sup>21</sup>His<sup>23</sup>Leu<sup>26</sup>Ala<sup>29</sup>Pro<sup>31</sup>Asp-h-amylin,  $^{13}$ Thr $^{18}$ Arg $^{21}$ His $^{23}$ Leu $^{28}$ . $^{19}$ Pro $^{31}$ Asp-h-amylin, and  $^{13} {\rm Thr}^{18} {\rm Arg}^{21} {\rm His}^{23} {\rm Leu}^{25} {\rm Pr} o^{26} {\rm Ala}^{28,29} {\rm Pro}^{31} {\rm Asp-h-amylin.}$ The method\according to any of claims 1-4,

wherein said amylin agonist is <sup>25,28,29</sup> Pro-h-amylin.

The method according to any of claims 1 or 2,

wherein said gastritis or gastric ulceration is associated

with the administration of a non-steroidal anti
inflammatory drug.

10. The method according to any of claims 3 or 4 wherein said non-steroidal anti-inflammatory agent is selected from the group consisting of salicylate, phenylbutazone, indomethacin, acetominophen, phenacetin, naproxen and ibúprofen.

11) A pharmaceutical composition comprising (1) an amylin or an amylin agonist, or a pharmaceutically acceptable salt thereof, wherein said amylin agonist is not

a calcitonin, and (2) a non-steroidal anti-inflammatory agent, in a pharmaceutically acceptable carrier and dose.

12. The pharmaceutical composition according to claim 11, wherein said non-steroidal anti-inflammatory agent is selected from the group consisting of salicylate, phenylbutazone, indomethacin, acetominophen, phenacetin, naproxen, and ibuprofen.

[PATENTS\224042\224042.APP]